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(71) Applicant (for all designated States except US): **ELI LILLY AND COMPANY** [US/US]; Lilly Corporate Center, Indianapolis, Indiana 46285 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **KOHLMAN, Daniel, Timothy** [US/US]; 6281 East Old Otto Court, Camby, Indiana 46113 (US). **VICTOR, Frantz** [US/US]; 4855 North Tuxedo Street, Indianapolis, Indiana 46205 (US). **XU, Yao-Chang** [US/US]; 10815 Timber Springs Drive East, Fishers, Indiana 46038 (US). **YING, Bai-Ping** [US/US]; 7717 Hidden Ridge, Fishers, Indiana 46038 (US). **ZHANG, Deyi** [US/US]; 1372 Kirklees Drive, Carmel, Indiana 46032 (US).

(74) Agents: **TUCKER, R., Craig** et al.; Eli Lilly and Company, P. O. Box 6288, Indianapolis, Indiana 46206-6288 (US).

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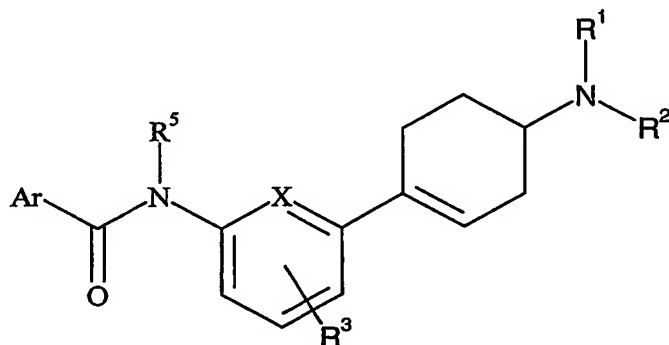
Declarations under Rule 4.17:

— as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii)) for the following designations AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, ARIPO patent (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)

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(54) Title: SUBSTITUTED (4-AMINOCYCLOHEXEN-1-YL)PHENYL AND (4-AMINOCYCLOHEXEN-1-YL)PYRIDINYL COMPOUNDS AS 5-HT_{1F} AGONISTS



(I)

(57) Abstract: The present invention relates to compounds of formula (I) or a pharmaceutically acceptable acid addition salt thereof, where; X is -C(R⁴)= or -N=; Ar is phenyl, substituted phenyl, heterocycle, or substituted heterocycle; R¹ and R² are independently hydrogen or C₁-C₃ alkyl; R³ is hydrogen, fluoro, or methyl; when X is -C(R⁴)=, R⁴ is hydrogen, fluoro, or methyl, provided that no more than one of R³ and R⁴ may be other than hydrogen; and R⁵ is hydrogen, methyl, or ethyl. The compounds of the present invention are useful for activating 5-HT_{1F} receptors, inhibiting dural

protein extravasation, and for the treatment or prevention of migraine in a mammal.